

CLAIMS:

1. A method of producing a library of mutant nucleic acid molecules comprising:
 - (a) obtaining a template nucleic acid;
 - 5 (b) preparing a first oligonucleotide corresponding to a first desired mutation within said template nucleic acid;
 - (c) preparing a second oligonucleotide corresponding to a second desired mutation within said template nucleic acid;
 - (d) mixing the oligonucleotides prepared in said steps (b) and (c) so as to
 - 10 hybridize said oligonucleotides to said template nucleic acid;
 - (e) subjecting the mixture of step (d) to the linear cyclic amplification reaction to produce a library of mutant template nucleic acids.
2. The method according to claim 1, wherein said oligonucleotides in said steps
- 15 (b) and (c) are discontinuous.
3. The method according to claim 1, wherein said step first and second oligonucleotides are present in less than saturation concentration.
- 20 4. The method according to claim 1, wherein the mixture of said step (d) further comprises non-mutagenic oligonucleotides corresponding to either or both of said first and second oligonucleotides.
5. The method according to claim 1, wherein said template nucleic acid
- 25 corresponds to a desired protein product.
6. The method according to claim 4, wherein said protein product comprises an enzyme, hormone, vaccine, peptide therapeutic or antibody.
- 30 7. The method according to claim 4, further comprising the steps of:
 - (f) transforming said mutant template nucleic acids from said library into a competent host cell;
 - (g) expressing protein corresponding to said mutant nucleic acids in said host cell;
 - 35 (h) screening said expressed proteins for desired characteristics.